(FILE 'HOME' ENTERED AT 12:34:19 ON 05 JAN 2011)

FILE 'MEDLINE, CAPLUS, BIOSIS, SCISEARCH, LIFESCI' ENTERED AT 12:34:59 ON 05 JAN 2011

- L1 34967 S (INDUC? OR ACTIVAT?) (5A) (TH1 OR HELPER(W)T)
- L2 179737 S TCR OR T(W)CELL(W)RECEPTOR
- L3 1912 S L1(P)L2
- L4 36213 S (PLASMID OR VECTOR OR POLYNUCLEOTIDE OR DNA OR NUCLEIC(W)ACID
- L5 126 S L1(P)L4
- L6 40 DUP REM L5 (86 DUPLICATES REMOVED)
- => d au ti so pi 20-40 16
- L6 ANSWER 20 OF 40 CAPLUS COPYRIGHT 2011 ACS on STN
- AU Hayakawa, Satoshi
- TI Human pregnancies and their complications from standpoints of Th1/Th2 balance and lymphocyte activation
- SO Nippon Sanka Fujinka Gakkai Zasshi (1999), 51(8), 626-632 CODEN: NISFAY; ISSN: 0300-9165
- L6 ANSWER 21 OF 40 MEDLINE on STN DUPLICATE 15
- AU Keino H; Matsumoto I; Okada S; Kurokawa M; Kato T; Tokuhisa T; Usui M; Taniquchi M; Nishioka K; Sumida T
- TI A single cell analysis of TCR AV24AJ18+ DN T cells.
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- L6 ANSWER 22 OF 40 MEDLINE on STN DUPLICATE 16
- AU Hill M; Beeson D; Moss P; Jacobson L; Bond A; Corlett L; Newsom-Davis J; Vincent A; Willcox N
- TI Early-onset myasthenia gravis: a recurring T-cell epitope in the adult-specific acetylcholine receptor epsilon subunit presented by the susceptibility allele HLA-DR52a.
- SO Annals of neurology, (1999 Feb) Vol. 45, No. 2, pp. 224-31. Journal code: 7707449. ISSN: 0364-5134. L-ISSN: 0364-5134.
- L6 ANSWER 23 OF 40 MEDLINE on STN DUPLICATE 17
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- TI Predominant expression of T helper 2 cytokines and altered expression of T helper 1 cytokines in long-term allograft survival induced by intrathymic immune modulation with donor class I major histocompatibility complex peptides.
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- AU Bourdette, Dennis N.; Chou, Yuan K.; Whitham, Ruth H.; Buckner, Jane; Kwon, Hi Jong; Nepom, Gerald T.; Buenafe, Abigail; Cooper, Shelley A.; Allegretta, Mark; Hashim, George A.; Offner, Halina; Vandenbark, Arthur A.
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- SO Journal of Immunology (1998), 161(2), 1034-1044 CODEN: JOIMA3; ISSN: 0022-1767
- L6 ANSWER 25 OF 40 MEDLINE on STN DUPLICATE 18
- AU Ariail K S; Bebo B F; Adlard K; Robey I; Burrows G; Newman M J; Todd C W; Vandenbark A A; Offner H
- TI Novel adjuvants for induction of T-cell and antibody responses to

- encephalitogenic and regulatory determinants in Lewis rats.
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- L6 ANSWER 26 OF 40 MEDLINE on STN DUPLICATE 19
- AU Zhang D H; Cohn L; Ray P; Bottomly K; Ray A
- TI Transcription factor GATA-3 is differentially expressed in murine Th1 and Th2 cells and controls Th2-specific expression of the interleukin-5 gene.
- SO The Journal of biological chemistry, (1997 Aug 22) Vol. 272, No. 34, pp. 21597-603.

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- TI Characterization of tumor-infiltrating T lymphocytes in B-cell lymphomas of mucosa-associated lymphoid tissue.
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- AU Ohara-Nemoto, Y.; Kaneko, M.
- TI Expression of T-cell receptor $V\beta2$ and type 1 helper T-cell-related cytokine mRNA in streptococcal pyrogenic exotoxin-C-activated human peripheral blood mononuclear cells
- SO Canadian Journal of Microbiology (1996), 42(11), 1104-1111 CODEN: CJMIAZ; ISSN: 0008-4166
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DUPLICATE 24

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- TI Borrelia burgdorferi activates a T helper type 1-like T cell subset in Lyme arthritis
- SO Journal of Experimental Medicine (1991), 174(3), 593-601 CODEN: JEMEAV; ISSN: 0022-1007
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- SO International Immunology (2003), 15(1), 1-10 CODEN: INIMEN; ISSN: 0953-8178
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- AU Lobito Adrian A; Yang Bingzhi; Lopes Marcela F; Miagkov Alexei; Adams Robert N; Palardy Gregory R; Johnson Michele M; McFarland Hugh I; Recher Michael; Drachman Daniel B; Lenardo Michael J
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- TI Immune tolerance induced by polyethylene glycol-conjugate of protein antigen: clonal deletion of antigen-specific Th-cells in the thymus.
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- => d ab 14 16
- L6 ANSWER 14 OF 40 CAPLUS COPYRIGHT 2011 ACS on STN DUPLICATE 10
- AΒ Interferon-regulatory factor-4 (IRF-4) is a member of the IRF family of transcription factors expressed in lymphocytes and macrophages. The previous studies using mice deficient in the IRF-4 gene showed profound defects in function of both B and T cells. To further investigate the role of IRF-4 in CD4+ T cell function, IRF-4-/- mice were challenged with the intracellular pathogen Leishmania major. The mice were protected against L. major during the early phase of the infection and CD4+ T cells of the infected mice produced IFN- $\!\gamma$ in response to L. major antigen. However, during the late phase of infection, lymphocyte nos. were dramatically reduced in the draining lymph nodes, resulting in the deterioration of the lesion, indicating that IRF-4 was required for sustained immune responses against L. major infection. The function of CD4+ T cells was further investigated using TCR transgenic mice lacking the IRF-4 gene. CD4+ T cells from IRF-4-/- mice produced IFN- γ and expressed T-bet after culture under Th1-skewing conditions in vitro. However, Th2 cell development was not observed after culture under $\operatorname{Th2-polarizing}$ conditions. Proliferation of $\operatorname{CD4+}$ T cells to ${
 m IL}{
 m -4}$ was reduced in ${
 m IRF}{
 m -4}{
 m -/-}$ mice, suggesting the defects in the responsiveness to IL-4. Furthermore, stimulation of the IRF-4-/- CD4+ T $\,$ cells with IL-4-induced activation of signal transducer and activator of transcription 6, but not expression of growth factor independent-1. Thus,

development of CD4+ T cell subsets differentially depends on IRF-4; induction of Th1 response does not depend on IRF-4, while Th2 response depends entirely on IRF-4.

=> d ab 29 16

L6 ANSWER 29 OF 40 CAPLUS COPYRIGHT 2011 ACS on STN

AΒ Alloreactive T cells recognize either determinants of the intact donor MHC mols. displayed on the surface of transplanted cells or peptide fragments of donor antigens associated with self-MHC mols. by their T cell receptors (TCR). To investigate the relation between the TCR β chain structure and allorecognition, the authors established and characterized four long-term T cell lines and seven T cell clones derived following a mixed lymphocyte reaction (MLR) between fully histoincompatible DA (RT1a) and LEW (RT11) rat lymph node cells. These DA anti-LEW T cells were phenotypically CD4+, CD8-, $\alpha\beta$ TCR+ and produced interferon- γ but not IL-4, consistent with being Th1 CD4+ T cells. As might be expected, these cells were not significantly cytotoxic and did not display suppressor activity. Anal. of the TCR β chain gene structure revealed a very restricted repertoire in both long-term lines and clones. The TCRBV6S1 gene was present in 15/21 of the alloreactive T cell mRNA transcripts but only 1/12 of unstimulated DA splenic TCR mRNA transcripts. Similarly, the TCRBJ2S1 gene was also used frequently in the alloreactive transcripts (17/21) but in only 2/12 unstimulated splenic transcripts. Furthermore, all 15 of the alloreactive TCRBV6S1 transcripts had a distinctive four amino acid N region motif not present in any of the unstimulated TCR transcripts. These expts. reveal a distinct homogeneity amongst stable allogeneic T cells in culture. If these results reflect the situation in vivo, the possibility exists that specific immunotherapy may be successful in preventing allograft rejection.